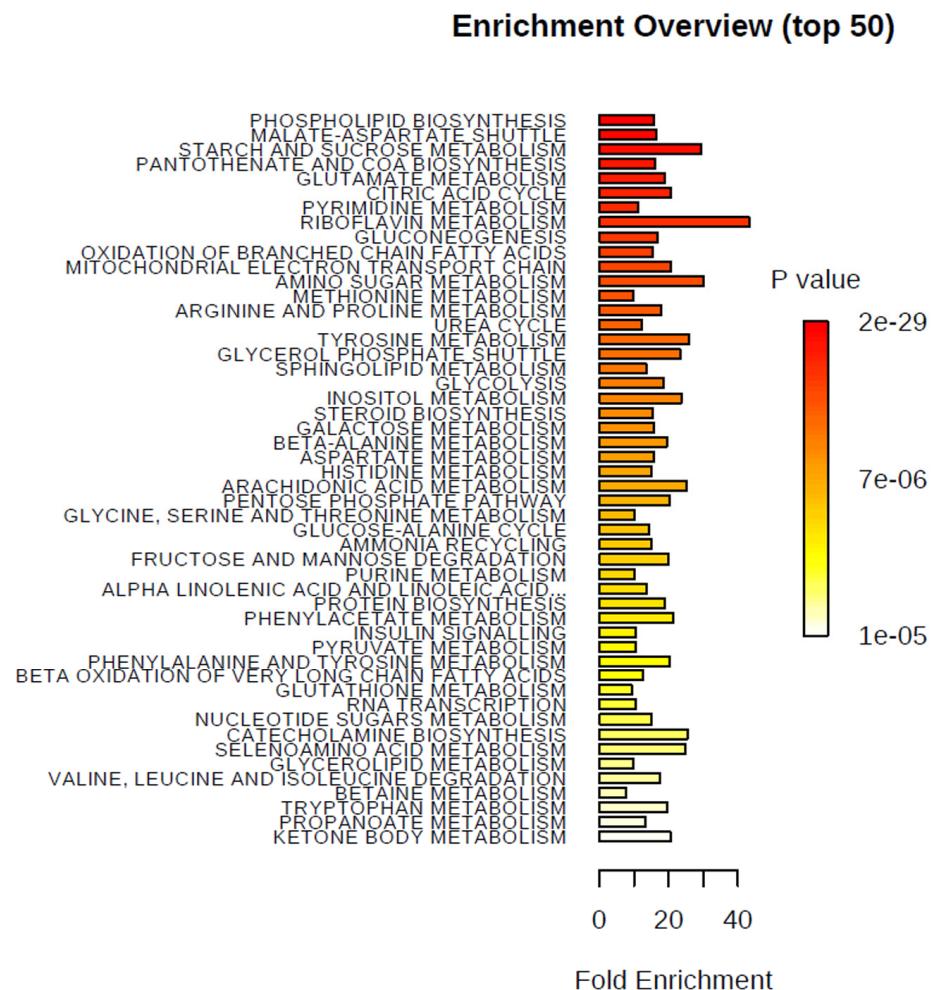
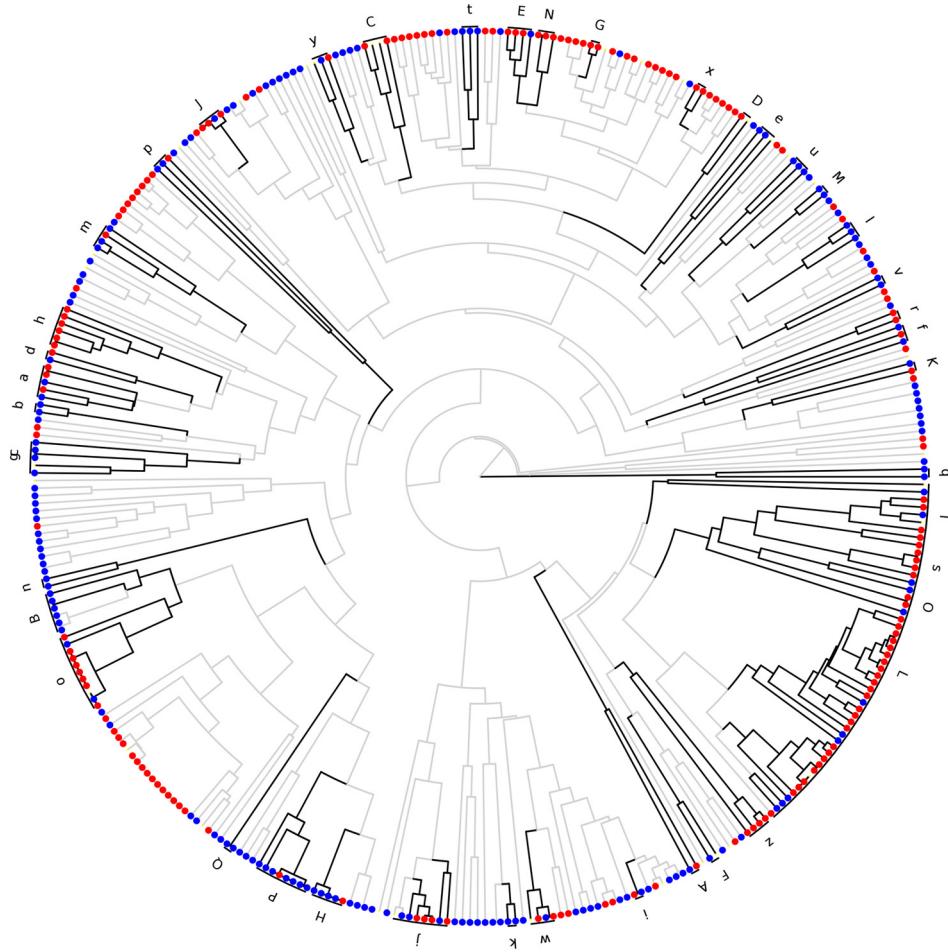


## SUPPLEMENTARY FIGURES

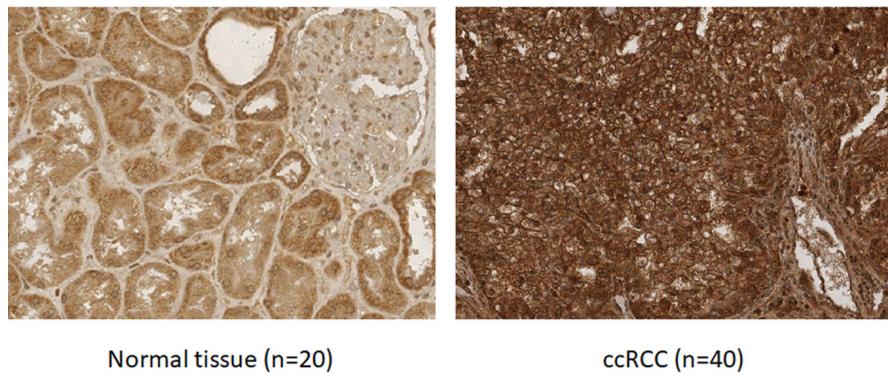


**Supplementary Figure 1.** Metabolite set enrichment overview performed in MetaboAnalyst 3.0

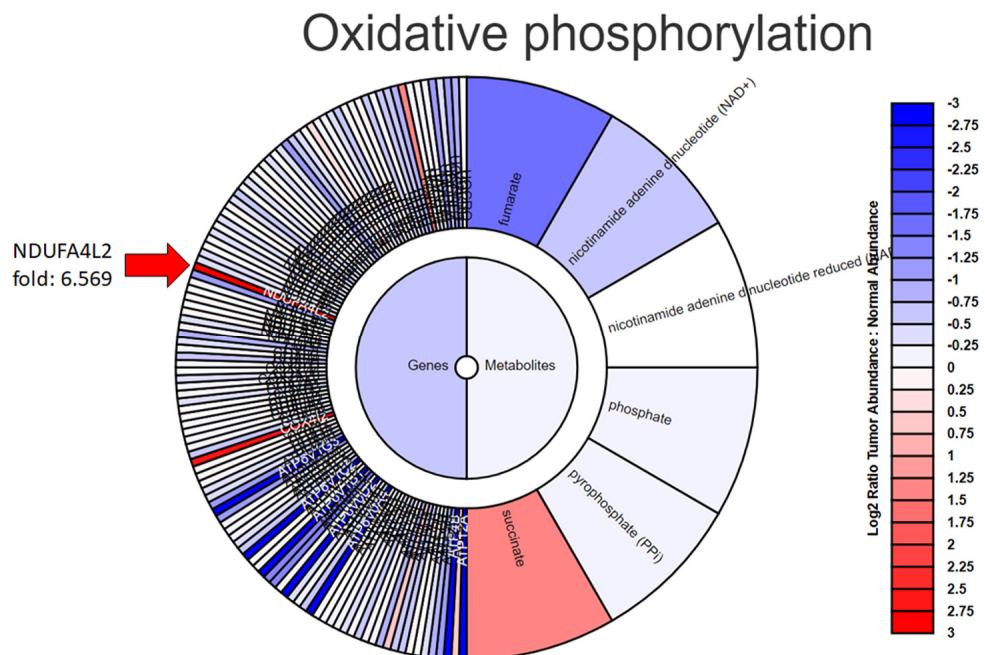


**Supplementary Figure 2.** Tanimoto chemical similarity mapping of all identified metabolites in ccRCC. Clusters are defined by comparing within- versus between group similarities, forming a clustered chemical similarity tree. Dark black lines indicate boundaries of clusters significantly different in tumor versus normal tissue ( $p < 0.05$ ). Cluster letter labels are detailed in Supplementary Table 2. Increased metabolites in ccRCC are labeled as red nodes. Decreased metabolites are blue nodes.

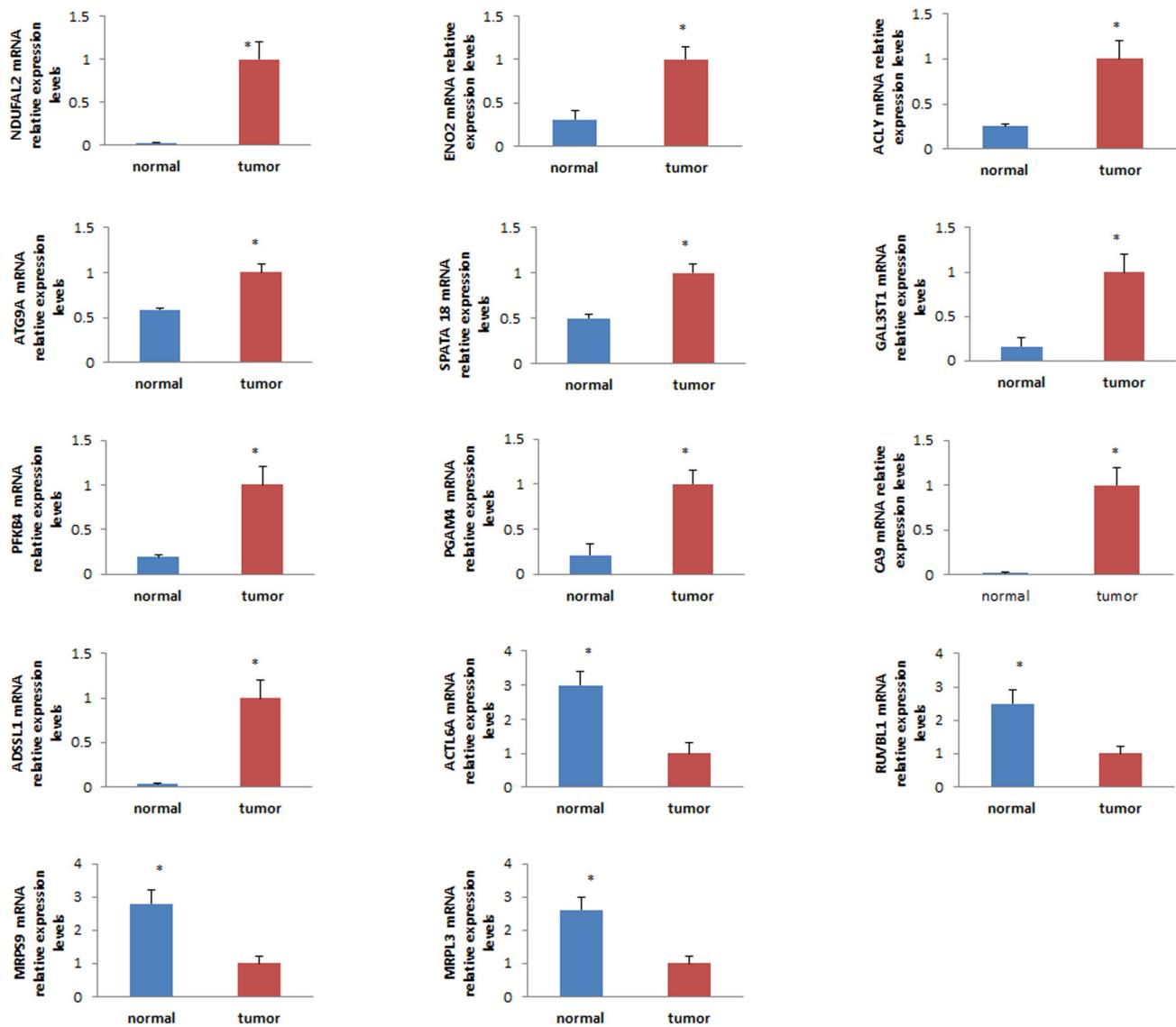
## GLUT 1 tissue expression



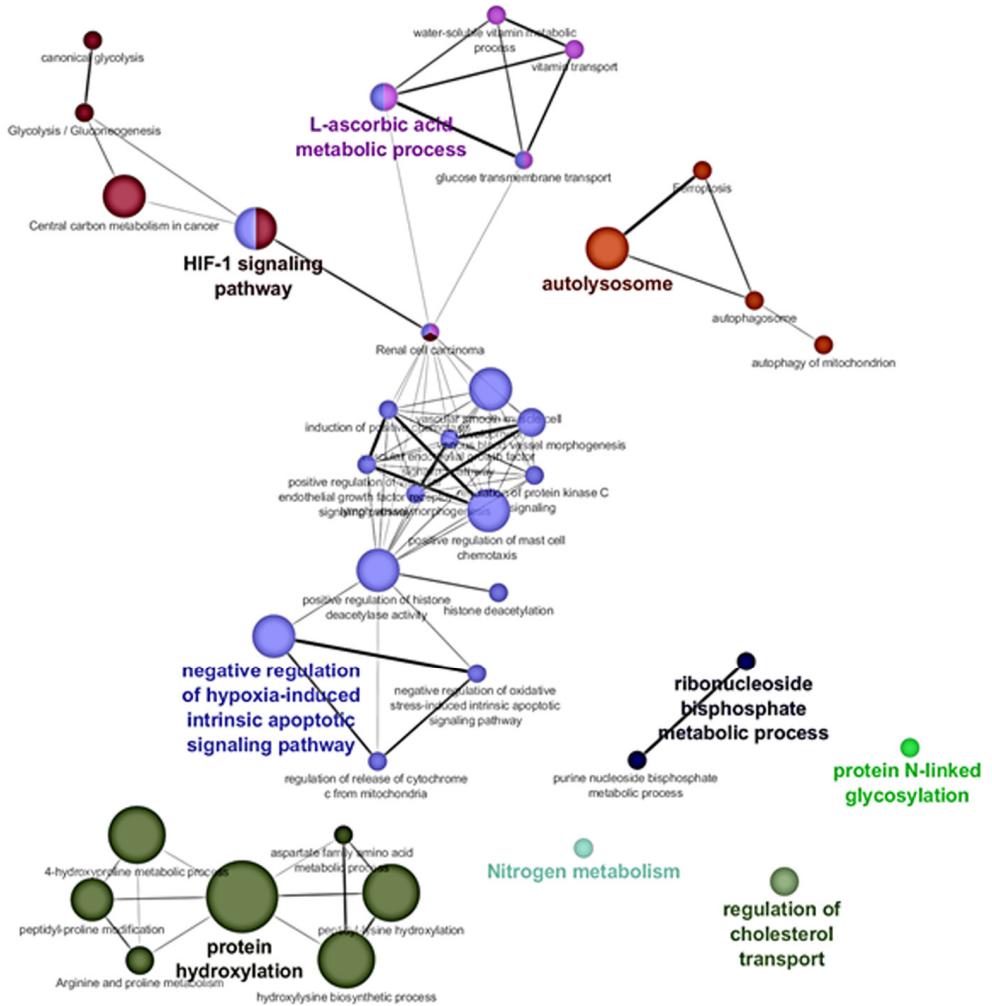
**Supplementary Figure 3.** Glut1 expression in normal (n=20) and ccRCC (n=40) specimens.



**Supplementary Figure 4.** Exploration of the “Metabogram” Data Portal. The changes in the “Oxidative phosphorylation” metabolism pathway are shown in both transcripts and metabolites when comparing tumors to adjacent normal kidney tissues.



**Supplementary Figure 5.** Gene expression by real-time PCR.



**Supplementary Figure 6.** Enriched GO network groups using ClueGO for GSE117890 dataset analysis. Biological processes (GO category) are visualized ( $\kappa$  score  $\geq 0.3$ ) as a functional grouped network and only the most significant interactions are shown. Each node represents a biological process. Edges represent connections between the nodes and the length of each edge reflects the relatedness of two processes.